



HBZ impedes the Menin function and up-regulates the transcription of the hTERT gene in leukemic cells

Malgorzata Borowiak, Anne-Sophie Kuhlmann, Sophie Girard, Louis Gazzolo, Madeleine Duc Dodon

► To cite this version:

Malgorzata Borowiak, Anne-Sophie Kuhlmann, Sophie Girard, Louis Gazzolo, Madeleine Duc Dodon. HBZ impedes the Menin function and up-regulates the transcription of the hTERT gene in leukemic cells. 15th International Conference on Human Retroviruses: HTLV and Related Viruses, Jun 2011, Leuven and Gembloux, Belgium. pp.A182, 10.1186/1742-4690-8-S1-A182 . inserm-00663652

HAL Id: inserm-00663652

<https://www.hal.inserm.fr/inserm-00663652>

Submitted on 27 Jan 2012

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



MEETING ABSTRACT

Open Access

HBZ impedes the Menin function and up-regulates the transcription of the hTERT gene in leukemic cells

Malgorzata Borowiak, Anne-Sophie Kuhlmann, Sophie Girard, Louis Gazzolo, Madeleine Duc Dodon*

From 15th International Conference on Human Retroviruses: HTLV and Related Viruses
Leuven and Gembloux, Belgium. 5-8 June 2011

Leukemic cells from Adult T-cell leukemia (ATL) patients display elevated telomerase activity, resulting mainly from transcriptional up-regulation of the human telomerase catalytic subunit (hTERT). We have previously shown that HBZ (HTLV-1 bZip) protein cooperates with JunD transcription factor to enhance hTERT expression after JunD anchoring to Sp1 bound to Sp1 sites within the hTERT proximal promoter. In normal somatic cells, telomerase expression is negatively regulated by tumor suppressor gene products, such as Menin, encoded by the multiple endocrine neoplasia type 1 (MEN-1) gene. Interestingly, the interaction of Menin with JunD has been shown to repress its transcriptional activity.

We report here that, in HBZ-expressing cells, Menin and HBZ exert opposite effects on JunD-mediated regulation of hTERT transcription. Chromatin immunoprecipitation as well as functional assays demonstrate that this antagonism is linked to the recruitment of p300 by HBZ and HDACs by Menin. Furthermore, knock-down of Menin in the ATL T1-Om1 (only expressing HBZ) cells results in an increase of hTERT expression and of telomerase activity, whereas a knock-down of HBZ exerts opposite effects. Interestingly, primary leukemic cells isolated from ATL patients that express high amounts of hTERT transcripts, are also characterized by an elevated expression of HBZ and MEN-1 genes. Thus, in leukemic cells, HBZ behaves as a key factor impeding the Menin function and sustaining hTERT transcription. These findings underline the critical role of HBZ as a

tumor-promoting protein during the development of the HTLV-1-induced leukemogenic process.

Published: 6 June 2011

doi:10.1186/1742-4690-8-S1-A182

Cite this article as: Borowiak et al.: HBZ impedes the Menin function and up-regulates the transcription of the hTERT gene in leukemic cells. *Retrovirology* 2011 **8**(Suppl 1):A182.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit



* Correspondence: mducdodo@ens-lyon.fr

Virologie Humaine, INSERM-U758, Ecole Normale Supérieure de Lyon, IFR
128 BioSciences Lyon-Gerland, Lyon, Cedex 07, 69364, France